

# Shao-Yan Hu

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/4017277/publications.pdf>

Version: 2024-02-01

6  
papers

29  
citations

2682572

2  
h-index

2272923

4  
g-index

7  
all docs

7  
docs citations

7  
times ranked

32  
citing authors

| # | ARTICLE   | IF   | CITATIONS |
|---|---|------|-----------|
| 1 | Ovarian germ cell tumor/mastocytosis with <i>KIT</i> mutation: A unique clinicopathological entity. <i>Genes Chromosomes and Cancer</i> , 2022, 61, 50-54.  | 2.8  | 2         |
| 2 | Dynamic change of variant allele frequency reveals disease status, clonal evolution and survival in pediatric relapsed B-cell acute lymphoblastic leukaemia. <i>Clinical and Translational Medicine</i> , 2022, 12, . | 4.0  | 1         |
| 3 | Arsenic Combined With All-Trans Retinoic Acid for Pediatric Acute Promyelocytic Leukemia: Report From the CCLG-APL2016 Protocol Study. <i>Journal of Clinical Oncology</i> , 2021, 39, 3161-3170.                     | 1.6  | 21        |
| 4 | Leukemic progenitor cells enable immunosuppression and post-chemotherapy relapse via IL-36-inflammatory monocyte axis. <i>Science Advances</i> , 2021, 7, eabg4167.   | 10.3 | 3         |
| 5 | Case Report: An Infant with Severe Thrombocytopenia Diagnosed with Type 2B von Willebrand Disease Due To a De Novo p.Val1316Met Mutation. <i>Turkish Journal of Haematology</i> , 2020, 37, 296-298.                  | 0.2  | 0         |
| 6 | Case Report: An Infant with Severe Thrombocytopenia Diagnosed with Type 2B von Willebrand Disease Due To a De Novo p.Val1316Met Mutation. <i>Turkish Journal of Haematology</i> , 2020, 37, 296-298.                  | 0.5  | 2         |